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## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification <sup>5</sup> :  C08F 10/00, 4/64	A1	(11) International Publication Number: <b>WO 93/19104</b>  (43) International Publication Date: 30 September 1993 (30.09.93)									
<p>(21) International Application Number: PCT/US93/02584</p> <p>(22) International Filing Date: 19 March 1993 (19.03.93)</p> <p>(30) Priority data:</p> <table border="0"><tr><td>07/857,886</td><td>26 March 1992 (26.03.92)</td><td>US</td></tr><tr><td>07/941,014</td><td>4 September 1992 (04.09.92)</td><td>US</td></tr><tr><td>08/008,003</td><td>21 January 1993 (21.01.93)</td><td>US</td></tr></table> <p>(60) Parent Application or Grant (63) Related by Continuation US 08/008,003 (C1P) Filed on 20 January 1993 (20.01.93)</p> <p>(71) Applicant (<i>for all designated States except US</i>): THE DOW CHEMICAL COMPANY [US/US]; 2030 Dow Center, Abbott Road, Midland, MI 48640 (US).</p>		07/857,886	26 March 1992 (26.03.92)	US	07/941,014	4 September 1992 (04.09.92)	US	08/008,003	21 January 1993 (21.01.93)	US	<p>(72) Inventors; and (75) Inventors/Applicants (<i>for US only</i>): ROSEN, Robert, K. [US/US]; 1800 Rivercrest Drive, No. 210, Sugar Land, TX 77478 (US). NICKIAS, Peter, N. [US/US]; 4512 North Saginaw Road, Apt. 1120, Midland, MI 48642 (US). DEVORE, David, D. [US/US]; 4881 East Shaffer Road, Midland, MI 48642 (US). STEVENS, James, C. [US/US]; 704 Georgetown Drive, Midland, MI 48642 (US). TIMMERS, Francis, J. [US/US]; 4605 Lund Drive, Midland, MI 48642 (US).</p> <p>(74) Agent: DELINE, Douglas, N.; The Dow Chemical Company, Patent Department, P.O. Box 1967, Midland, MI 48641-1967 (US).</p> <p>(81) Designated States: CA, FI, JP, KR, US, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).</p> <p>Published <i>With international search report.</i></p>
07/857,886	26 March 1992 (26.03.92)	US									
07/941,014	4 September 1992 (04.09.92)	US									
08/008,003	21 January 1993 (21.01.93)	US									
<p>(54) Title: ADDITION POLYMERIZATION CATALYSTS COMPRISING REDUCED OXIDATION STATE METAL COMPLEXES</p> <p>(57) Abstract</p> <p>Addition polymerization catalysts comprising a metal complex containing a Group 4 metal in the +3 oxidation state, a delocalized <math>\pi</math>-bonding moiety, and at least one stabilizing ligand, optionally in combination with an activating cocatalyst.</p>											

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ADDITION POLYMERIZATION CATALYSTS COMPRISING  
REDUCED OXIDATION STATE METAL COMPLEXES

This invention relates to addition polymerization catalysts comprising certain  
5 stabilized, reduced metal complexes. In one embodiment such catalysts additionally employ an  
activating cocatalyst. In another embodiment such catalysts may be employed with or without  
an activating cocatalyst. Finally the invention relates to an improved method for preparing the  
stabilized reduced metal complexes. The compositions are especially useful as catalysts for the  
polymerization of olefins such as ethylene for the preparation of polymers having utility as  
10 molding resins and in the formation of foamed products having cushioning and insulating  
applications and films that are useful as protective wrappings.

In U.S. Serial No. 545,403, filed July 3, 1990 (equivalent to EP-A-0416,845) there  
are disclosed and claimed certain monocyclopentadienyl metal complexes having utility as  
homogeneous olefin polymerization catalysts. In US-A-5,064,802 (equivalent to  
15 EP-A-0418,044), cationic monocyclopentadienyl metal complexes with salts of Bronsted acids  
containing a non-coordinating compatible anion are disclosed and claimed. In U.S. Serial No.  
547,718, also filed on July 3, 1990 (EP-A-0468,651), an oxidative activation technique for  
preparing such cationic catalysts is disclosed and claimed.

In US-A-4,057,565 there are disclosed Ti, Zr, or Hf derivatives of 2-dialkylamino-  
20 benzyl or 2-dialkylaminomethylphenyl all in the + 4 oxidation state which are useful as  
components of catalysts for olefin polymerization.

In J. Am. Chem. Soc. 100, 8068-8073 (1978) there is mentioned the synthesis and  
characterization of Ti(+3) complexes containing cyclopentadienyl groups and 2-dialkylamino-  
benzyl or 2-dialkylaminomethylphenyl groups. No mention of utility as addition  
25 polymerization catalysts is given.

In US patent 4,870,042, catalysts for olefin polymerizations comprising a pyrazolyl  
borate complex of titanium or zirconium compounds including titanium trichloride (Example 4)  
are disclosed.

In Organometallics, 10, 3227-3237 (1991) certain titanium + 3 complexes  
30 containing cyclopentadienyl groups and alkyl groups are mentioned. On page 3236, the  
reference states:

"So far no well-defined neutral titanium-based molecular system with established  
activity for catalytic olefin polymerization has been described... Apparently, the  
tervalent  $Cp^*_2TiR$  system cannot induce sufficient positive charge at the  $\beta$ -carbon  
atom of an incoming ethylene molecule to reach the polar transition state for  
35 migratory insertion."

In Journal of Organometallic Chemistry, 334 (1987)  $C1-C4, \eta^3\text{-allyl(bis-}\eta^5\text{-}$   
cyclopentadienyl)titanium (III) activated with dimethylaluminum chloride was found to create

a coordinated species  $\text{Cp}_2\text{Ti}(\text{allyl})-(\text{CH}_3)_2\text{AlCl}$  which was found to polymerize ethylene to a small extent. Use of stronger Lewis acids so as to cause ligand abstraction thereby destroying the requisite carbon-titanium bond was to be avoided according to the reference.

Figure 1 is an electron spin resonance spectrogram of the product obtained in preparation ii) of Examples 1-58.

Figure 2 is an electron spin resonance spectrogram of the product obtained in Example 59.

Figure 3 is an electron spin resonance spectrogram of the product obtained in Example 60.

According to the present invention in its most generic description, there is provided an addition polymerization catalyst comprising:

(i) a metal complex corresponding to the formula:  $\text{Cp}_a(\text{ZY})_b\text{ML}_c$ , wherein:

a is 1 or 2;

b is 0 or 1;

c is 1 or 2;

the sum of a, b and c is 3;

Cp independently each occurrence is a cyclopentadienyl group  $\pi$ -bound to M, or a hydrocarbyl, silyl, halo, halohydrocarbyl, hydrocarbylmetaloid or halohydrocarbylmetaloid substituted derivative of said cyclopentadienyl group, said Cp containing up to 50 nonhydrogen atoms, and, when a is 2, optionally both Cp groups may be joined together by a bridging group;

L independently each occurrence is hydride, halo, or a monovalent anionic ligand selected from covalently bonded hydrocarbyl, silyl, amido, phosphido, alkoxy, aryloxy, and sulfido groups optionally being further substituted with amine, phosphine, ether, and thioether; mixtures thereof; said ligand having up to 50 nonhydrogen atoms, with the proviso that in at least one occurrence L is a stabilizing ligand comprising an amine, phosphine, ether or thioether functionality able to form a coordinate-covalent bond or chelating bond with M, or except when a is 1, comprising an ethylenic unsaturation able to form an  $\eta^3$  bond with M;

M is a metal of Group 4 of the Periodic Table of the Elements in the +3 oxidation state;

Y is a linking group comprising nitrogen, phosphorus, oxygen or sulfur covalently bonded to M and Z through said nitrogen, phosphorus, oxygen or sulfur atom;

Z is a divalent moiety comprising oxygen, nitrogen, phosphorous, boron, or a member of Group 14 of the Periodic Table of the Elements, said moiety having up to 30 nonhydrogen atoms;

the ligand moiety consisting of -Cp-Z-Y- being dianionic and having the ionic charges residing formally on Cp and Y;

and, when b is 0 and optionally when b is 1,

(ii) an activating cocatalyst.

In accordance with one embodiment of the above generic description of the invention there is provided an addition polymerization catalyst comprising in combination:

(i) a metal complex, A<sub>1</sub>, corresponding to the formula:



wherein:

Cp' independently each occurrence is a cyclopentadienyl group  $\eta$ -bound to M, or a hydrocarbyl, silyl, halo, halohydrocarbyl, hydrocarbylmetaloid or halohydrocarbylmetaloid substituted derivative of said cyclopentadienyl group, said Cp' containing up to 50  
10 nonhydrogen atoms, and optionally both Cp groups may be joined together by a bridging group;

M is a metal of Group 4 of the Periodic Table of the Elements in the +3 oxidation state;

L' is a monovalent anionic stabilizing ligand selected from the group consisting  
15 of: covalently bonded hydrocarbyl, silyl, amido, phosphido, alkoxy, aryloxy, sulfido groups and mixtures thereof, said group being further substituted with an amine, phosphine, ether, or thioether containing substituent able to form a coordinate-covalent bond or chelating bond with M; said ligand having up to 50 nonhydrogen atoms; and  
(ii) an activating cocatalyst.

20 Preferably in accordance with the present invention the ratio of metal complex to activating cocatalyst is from 1:0.01 to 1:10<sup>6</sup>.

In a further embodiment there is provided an addition polymerization catalyst comprising in combination:

(i) a metal complex, A<sub>2</sub>, corresponding to the formula



wherein:

Cp'' is a cyclopentadienyl group  $\eta$ -bonded to M, or a hydrocarbyl, silyl, halo, halohydrocarbyl, hydrocarbylmetaloid, or halohydrocarbylmetaloid substituted derivative thereof, said Cp'' containing up to 50 nonhydrogen atoms;

30 M is a metal of Group 4 of the Periodic Table of the Elements in the +3 oxidation state;

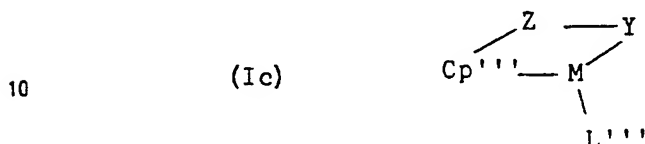
L'' independently each occurrence is hydride, halo, or a monovalent anionic ligand selected from the group consisting of covalently bonded hydrocarbyl, silyl, amido, phosphido, alkoxy, aryloxy, and sulfido groups; mixtures thereof; and amine, phosphine,  
35 ether, and thioether derivatives of the foregoing, said ligand having up to 50 nonhydrogen atoms, with the proviso that in at least one occurrence L'' is a stabilizing ligand comprising an amine, phosphine, ether or thioether functionality able to form a coordinate-covalent bond or

chelating bond with M, or comprising an ethylenic unsaturation able to form an  $\eta^3$  bond with M; and

(ii) an activating cocatalyst.

Preferably the activating cocatalyst is present in an amount to provide a ratio of  
5 metal complex to activating cocatalyst from 1:0.01 to 1:10<sup>6</sup>.

In a still further embodiment of the present invention there is provided a metal complex, A<sub>3</sub>, corresponding to the formula:



wherein:

M is a metal of Group 4 of the Periodic Table of the Elements in the +3 oxidation  
15 state;

Cp''' is a cyclopentadienyl group, or a hydrocarbyl, silyl, halo, halohydrocarbyl, or hydrocarbylmetalloid substituted derivative thereof, said Cp''' containing up to 50 nonhydrogen atoms;

Z is a divalent moiety comprising oxygen, nitrogen, phosphorous, boron, or a  
20 member of Group 14 of the Periodic Table of the Elements said moiety having up to 30 nonhydrogen atoms;

Y is a linking group comprising nitrogen, phosphorus, oxygen or sulfur covalently bonded to M and Z through said nitrogen, phosphorus, oxygen or sulfur atom, the ligand moiety consisting of -Cp'''-Z-Y- being dianionic and having the ionic charges residing formally  
25 on Cp''' and Y; and

L''' is a monovalent stabilizing ligand selected from the group consisting of L' and C<sub>3-15</sub> hydrocarbyl groups comprising an ethylenic unsaturation able to form an  $\eta^3$  bond with M.

The above complexes, A<sub>3</sub>, are suitable for use in polymerization of addition  
30 polymerizable monomers alone or optionally in the presence of an activating cocatalyst. In the latter event there is provided an addition polymerization catalyst comprising:

- (i) the complex, A<sub>3</sub>, and
- (ii) an activating cocatalyst.

The activating cocatalyst is preferably used in an amount to provide a ratio of  
35 metal complex to activating cocatalyst from 1:0.01 to 1:10<sup>6</sup>.

It is believed, without agreeing to be bound by such belief, that the metal complexes A<sub>1</sub>, A<sub>2</sub>, and A<sub>3</sub> when combined with the activating cocatalyst are converted into

cationic metal complexes wherein the metal is in the + 3 oxidation state. Evidence for this belief is established by electron spin resonance spectroscopy (ESR) and by the fact that solutions of the metal complexes in combination with the activating cocatalyst, especially tris(perfluorophenyl)boron are conductive in cyclic voltammetry analysis without added supporting electrolyte, whereas the complex alone is not conductive under the same conditions.

The presence of the above identified stabilizing ligand in the complexes of the invention results in improved complex stability and greatly improved usefulness in the formation of catalysts. The catalysts formed from these complexes by combination with the activating cocatalyst additionally possess higher activity and are more efficient addition polymerization catalysts than catalysts formed from corresponding complexes lacking such stabilizing ligand group.

The metal complexes, A<sub>1</sub>, may be prepared by a process comprising contacting a Group 4 metal complex corresponding to the formula:



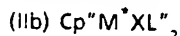
wherein Cp' and L' are as previously defined, and

M\* is a metal of Group 4 of the Periodic Table of the Elements in the +4 oxidation state; and

X is halide or C<sub>1-10</sub> hydrocarbyloxy,

with a reducing agent under reducing conditions to form the Group 4 metal complex.

The metal complexes, A<sub>2</sub>, may be prepared by a process comprising contacting a Group 4 metal complex corresponding to the formula:



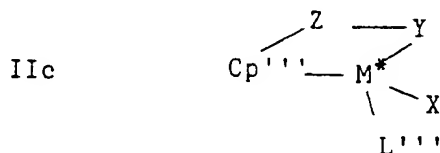
wherein Cp'' and L'' are as previously defined, and

M\* is a metal of Group 4 of the Periodic Table of the Elements in the +4 oxidation state; and

X is halide or C<sub>1-10</sub> hydrocarbyloxy,

with a reducing agent under reducing conditions to form the Group 4 metal complex.

Finally, the metal complexes, A<sub>3</sub>, may be prepared by a process comprising contacting a Group 4 metal complex corresponding to the formula:



wherein Cp''', Z, Y, M\*, X and L''' are as previously defined,

with a reducing agent under reducing conditions to form the Group 4 metal complex.

There are also provided polymerization processes comprising contacting an addition polymerizable monomer or monomers with the foregoing catalysts under polymerization conditions and recovering the resulting polymer.

All reference to the Periodic Table of the Elements herein shall refer to the  
5 Periodic Table of the Elements, published and copyrighted by CRC Press, Inc., 1989. Also, any reference to a Group or Groups shall be to the Group or Groups as reflected in this Periodic Table of the Elements using the IUPAC system for numbering groups.

By the term "stabilizing ligand" is meant that the ligand group stabilizes the metal complex through either:

- 10           1) a nitrogen, phosphorus, oxygen or sulfur chelating bond, or  
            2) an  $\eta^3$  bond with a resonant, delocalized  $\pi$ -electronic structure.

Examples of stabilizing ligands of group 1) for use according to the present invention include alkyl, cycloalkyl, aryl, silyl, amido or phosphido ligands substituted with one or more aliphatic or aromatic ether-, thioether-, amine- or phosphine- functional groups,  
15 especially such amine or phosphine groups that are tertiary substituted, said stabilizing ligand having from 3 to 30 nonhydrogen atoms. Most preferred group 1) stabilizing ligands are 2-dialkylaminobenzyl or 2-(dialkylaminomethyl)phenyl groups containing from 1 to 4 carbons in the alkyl groups.

Examples of stabilizing ligands of group 2) for use according to the present invention, include linear  $C_{3-15}$  hydrocarbyl groups containing ethylenic unsaturation, such as  
20 allyl, 1-methylallyl, 2-methylallyl, 1,1-dimethylallyl, 1,2,3-trimethylallyl, 1-phenyl-3-benzylallyl or 1,1-diphenyl-3-(diphenylmethyl)allyl groups.

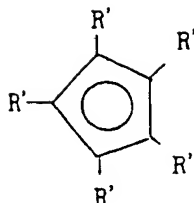
Preferred substituents of the groups Cp, Cp" and Cp''' are hydrocarbyl (including hydrocarbylene) and halosubstituted hydrocarbyl groups, said groups having from 1 to 30  
25 carbons, and  $C_{1-30}$  hydrocarbyl or halo-hydrocarbyl substituted metalloid radicals wherein the metalloid is an element from Group 14 of the Periodic Table of the Elements.

Exemplary hydrocarbyl radicals include straight and branched alkyl radicals, cyclic aliphatic hydrocarbon radicals, alkyl-substituted cyclic aliphatic hydrocarbon radicals, aromatic radicals and alkyl-substituted aromatic radicals. Preferred are methyl, ethyl, butyl and phenyl  
30 radicals. Exemplary organometalloid radicals include straight and branched chain silyl radicals, alkyl-substituted silyl radicals, germyl radicals and divalent derivatives of the foregoing. Preferred are trimethylsilyl, triethylsilyl, ethyldimethylsilyl, methyldiethylsilyl, dimethyl-t-butylsilyl, triphenylsilyl, triphenylgermyl, and trimethylgermyl radicals.

More particularly, suitable cyclopentadienyl or substituted cyclopentadienyl  
35 groups in complexes  $A_1$  are illustrated by the formula:



(III)



wherein:

R' is hydrogen, or a group selected from silyl, hydrocarbyl, and combinations thereof having up to 30 carbon or silicon atoms, or two R' groups together form a divalent derivative of such group.

Preferably, R' independently each occurrence is hydrogen, methyl, ethyl, propyl, butyl, pentyl, hexyl, (including where appropriate all isomers), cyclopentyl, cyclohexyl, norbornyl, benzyl, or phenyl or two R' groups are linked together thereby forming an indenyl, tetrahydroindenyl, fluorenyl, tetrahydrofluorenyl, or octahydrofluorenyl group in place of the cyclopentadienyl group.

Exemplary metal complexes of formula A<sub>1</sub> are: biscyclopentadienyl 2-(N,N-dimethylamino)benzyl titanium, bis(methylcyclopentadienyl) 2-(N,N-dimethylamino)benzyl titanium, bis(t-butylcyclopentadienyl) 2-((N,N-dimethylamino)methyl)phenyl titanium, bis(pentamethylcyclopentadienyl) 2-(N,N-dimethylamino)benzyl titanium, bis(indenyl)(2-N,N-dimethylamino)benzyl titanium, bis(fluorenyl)(2-N,N-dimethylamino)benzyl titanium, and corresponding zirconium and hafnium derivatives.

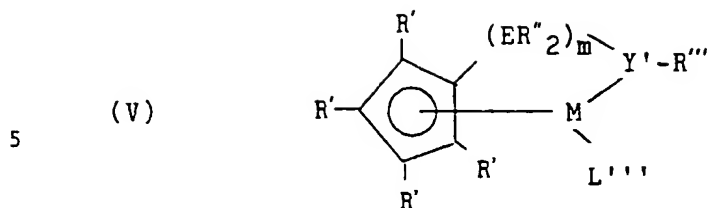
Additional bis-cyclopentadienyl compounds of formula A<sub>1</sub> include those containing a bridging group linking the cyclopentadienyl groups. Preferred bridging groups are those corresponding to the formula (ER''<sub>2</sub>)<sub>x</sub> wherein E is silicon or carbon, R'', independently each occurrence is hydrogen or a group selected from silyl, hydrocarbyl and combinations thereof, said R'' having up to 30 carbon or silicon atoms, and x is 1 to 8. Preferably R'' independently each occurrence is methyl, benzyl, tert-butyl, or phenyl.

Examples of the foregoing bridged cyclopentadienyl group containing complexes are compounds corresponding to the formula:

Exemplary complexes of formula  $A_2$  include cyclopentadienyl titanium allyl chloride, t-butylcyclopentadienyl titanium allyl chloride, pentamethylcyclopentadienyl titanium allyl chloride, pentamethylcyclopentadienyl titanium allyl methoxide, n-butylcyclopentadienyl titanium allyl isopropoxide, pentamethylcyclopentadienyl titanium allyl phenoxide, cyclopentadienyl titanium (2-methylallyl) chloride, ethylcyclopentadienyl titanium (2-methylallyl) ethoxide, cyclopentadienyl titanium 2-(N,N-dimethylamino)benzyl chloride, pentamethylcyclopentadienyl titanium 2-(N,N-dimethylamino)benzyl bromide, cyclopentadienyl titanium 2-(N,N-dimethylamino)benzyl methoxide, n-butylcyclopentadienyl titanium 2-((N,N-dimethylamino)methyl)phenyl isopropoxide, cyclopentadienyl titanium allyl dimethylamide, n-butylcyclopentadienyl titanium allyl dimethylamide, t-butylcyclopentadienyl titanium allyl diethylamide, cyclopentadienyl titanium diallyl, cyclopentadienyl titanium 2-(methylallyl) dimethylamide, cyclopentadienyl titanium (2-methylallyl) (di-t-butylamide), cyclopentadienyl titanium 2-(N,N-dimethylamino)benzyl dimethylamide, pentamethylcyclopentadienyl titanium 2-(N,N-dimethylamino)benzyl dimethylamide, methylcyclopentadienyl titanium methyl 2-(N,N-dimethylaminobenzyl), methylcyclopentadienyl titanium benzyl 2-(N,N-dimethylamino)benzyl, methylcyclopentadienyl titanium trimethylsilylmethyl 2-(N,N-dimethylamino)benzyl, pentamethylcyclopentadienyl titanium methyl 2-(N,N-dimethylaminobenzyl), pentamethylcyclopentadienyl titanium trimethylsilylmethyl 2-(N,N-dimethylamino)benzyl, and corresponding zirconium and hafnium complexes.

Illustrative cyclic complexes,  $A_3$ , that may be employed in the practice of the present invention include: (tert-butylamido)dimethyl(tetramethyl- $\eta^5$ -cyclopentadienyl)silane 2-(N,N-dimethylamino)benzyl titanium, (phenylamido)dimethyl(tetramethyl- $\eta^5$ -cyclopentadienyl)silane 2-(N,N-dimethylamino)benzyl titanium, (tert-butylamido)(tetramethyl- $\eta^5$ -cyclopentadienyl)-1,2-ethanediyl allyl titanium, (tert-butylamido)(tetramethyl- $\eta^5$ -cyclopentadienyl)-1,2-ethanediyl 2-(dimethylamino)benzyltitanium, (methylamido)(tetramethyl- $\eta^5$ -cyclopentadienyl)-1,2-ethanediyl allyl titanium, (tert-butylamido)dibenzyl-(tetramethyl- $\eta^5$ -cyclopentadienyl)silane 2-(N,N-dimethylamino)benzyl titanium, (benzylamido)dimethyl( $\eta^5$ -cyclopentadienyl)silane 2-((N,N-dimethylamino)methyl)phenyl titanium, (phenylphosphido)dimethyl(tetramethyl- $\eta^5$ -cyclopentadienyl)silane 2-(dimethylamino)benzyl titanium, (tert-butylamido)-dimethyl(tetramethyl- $\eta^5$ -cyclopentadienyl)silane allyl titanium, (methylamido)cyclotetramethylene( $\eta^5$ -octahydrofluorenyl)silane 2-(N,N-dimethylamino)benzyl titanium, (methylamido)cyclotetramethylene( $\eta^5$ -octahydrofluorenyl)silane 2-(N,N-dimethylamino)benzyl titanium, and corresponding zirconium or hafnium complexes.

Preferred cyclic complexes,  $A_3$ , are amidosilane- or amidoalkanediyl- compounds corresponding to the formula:



wherein:

M, L'', E, R', and R'' are as previously defined;

Y' is nitrogen;

R''' is a group selected from silyl, hydrocarbyl and combinations thereof, said group or combination having up to 30 carbon or silicon atoms; and

m is 1 or 2.

Preferably R''' is methyl, ethyl, propyl, butyl, pentyl, hexyl, (including where appropriate all isomers), cyclopentyl, cyclohexyl, norbornyl, benzyl, or phenyl.

In the most preferred embodiment  $-(ER''_2)_m-Y'R'''$  is an amidosilane or amidoalkane group of up to 30 nonhydrogen atoms, especially, (tert-butyl-amido)(dimethylsilyl), (methanido)(dimethylsilyl), or (tert-butylamido)-1-ethane-2-yl.

Other complexes which are useful in the catalyst compositions of this invention will be apparent to those skilled in the art.

Highly preferably in complexes A<sub>1</sub>, M is titanium and L' is 2-dimethylaminobenzyl.

Highly preferably in complexes A<sub>2</sub>, M is titanium and L'' is 2-dimethylaminobenzyl or allyl.

Highly preferably in complexes A<sub>3</sub>, M is titanium and L''' is 2-dimethylaminobenzyl or allyl.

By the term "reducing agent" herein is meant a metal or compound which, under reducing conditions causes M to be reduced from the + 4 oxidation state to the + 3 oxidation state. Examples of suitable metal reducing agents are alkali metals, alkaline earth metals, lead and zinc, alloys of alkali metals or alkaline earth metals such as sodium/mercury amalgam and sodium/potassium alloy. Examples of suitable reducing agent compounds are sodium naphthalenide, potassium graphite, and Grignard reagents. Most preferred reducing agents are the alkali metals or alkaline earth metals, especially magnesium.

By the term "reducing conditions" is meant the use of diluents and temperatures that allow the desired reduction to take place. Preferred temperatures are from 0°C to 200°C, more preferably from 5°C to 120°C and most preferably from 10°C to 50°C. Preferred diluents are polar solvents, more preferably C<sub>1-6</sub> aliphatic ethers, most preferably tetrahydrofuran, 1,2-dimethoxyethane or diethyl ether.

The term "activating cocatalyst" as used herein refers to a secondary component that renders the metal complex catalytically effective or improves the catalytic effectiveness of

the metal complex as an addition polymerization catalyst. Examples of the activating cocatalysts for use herein include aluminum compounds containing an Al-O bond such as the alkylalumoxanes, especially methylalumoxane and isobutyl modified methylalumoxane; aluminum alkyls; aluminum halides; alkylaluminum halides; Lewis acids other than any of the foregoing list; and mixtures of the foregoing.

Preferred Lewis acids are those compounds corresponding to the formula:  $R''''_3B$ , wherein  $R''''$  independently each occurrence is selected from hydrogen, silyl, hydrocarbyl, halohydrocarbyl, alkoxide, aryloxy, amide or combinations thereof, said  $R''''$  having up to 30 nonhydrogen atoms.

It is to be appreciated by those skilled in the art, that the above formula for the preferred Lewis acids represents an empirical formula, and that many Lewis acids exist as dimers or higher oligomers in solution or in the solid state. Other Lewis acids which are useful in the catalyst compositions of this invention will be apparent to those skilled in the art.

Preferred activating cocatalysts include trimethylaluminum, triisobutylaluminum, methylalumoxane, ethylalumoxane, chlorodiethylaluminum, dichloroethylaluminum, triethylboron, trimethylboron, triphenylboron and halogenated, especially fluorinated, triphenyl boron compounds.

Most highly preferred activating cocatalysts include triethylaluminum, methylalumoxane, and fluoro-substituted triaryl borons such as tris(4-fluorophenyl)boron, tris(2,4-difluorophenyl)boron, tris(3,5-bis(trifluoromethyl)phenyl)boron, tris(pentafluorophenyl)boron, pentafluorophenyldiphenylboron, and bis(pentafluorophenyl)phenylboron. Such fluoro-substituted triarylboranes may be readily synthesized according to techniques such as those disclosed in Marks, et al. J. Am. Chem. Soc., 113, 3623-3625 (1991).

A neutral Lewis base may additionally be present in combination with the complexes and catalysts of the invention. Suitable neutral Lewis bases include, for example, organic compounds containing oxygen, nitrogen, phosphorus or sulfur, or olefins. More specifically, examples of suitable electron donating compounds include organic amines, amides, ketones, nitriles, phosphines, phosphorylamides, esters, ethers (including cyclic ethers such as furans), thioethers and thioesters. Preferred are such compounds containing up to 20 nonhydrogen atoms. A highly preferred electron donating agent is tetrahydrofuran (THF).

It has been previously reported that Lewis bases such as THF have a severe degrading effect on the catalytic activity of cyclopentadienyl complexes of Group 4 metals in the +4 oxidation state. Jordan, et al., J. Am. Chem. Soc., 108, 7410-7411 (1986) at 7410, reported that "Addition of THF or other donor ligands slows the polymerization rate dramatically and in THF or  $CH_3CN$  solvents no activity is observed". Similarly, Bochmann et al., Polyhedron, 8 (13-14), 1838-1843 (1989) reported that bis(pentamethylcyclopentadienyl)titanium(IV) methyl tetraphenylborate prepared in THF solvent was inactive for ethylene

polymerization. Surprisingly, the presence of a neutral Lewis base does not necessarily adversely affect the catalytic activity of the metal complexes of the present invention, particularly the complexes of formula A<sub>3</sub>.

The catalyst can be formed by combining the activating cocatalyst (where  
5 required) with the complex, with or without a neutral Lewis base, optionally in the presence of a diluent. Complexes that need no activating cocatalyst may be used to polymerize addition polymerizable monomers without further additives or in combination with the above neutral Lewis base and optionally in the presence of a diluent. The preparation may be conducted in the presence of one or more addition polymerizable monomers, if desired. Preferably, the  
10 catalysts are prepared at a temperature within the range from -100°C to 300°C, preferably 0°C to 250°C, and most preferably 0°C to 100°C. Suitable solvents include straight and branched-chain hydrocarbons such as isobutane, butane, pentane, hexane, heptane, octane, and mixtures thereof; cyclic and alicyclic hydrocarbons such as cyclohexane, cycloheptane, methylcyclohexane, methylcycloheptane; perfluorinated hydrocarbons such as perfluorinated  
15 C<sub>4-10</sub> alkanes; and aromatic and alkyl-substituted aromatic compounds such as benzene, toluene and xylene. Suitable solvents also include liquid olefins which may act as monomers or comonomers including ethylene, propylene, butadiene, cyclopentene, 1-hexene, 3-methyl-1-pentene, 4-methyl-1-pentene, 1,4-hexadiene, 1-octene, 1-decene, styrene, divinylbenzene, 4-vinylcyclohexene, allylbenzene and vinyltoluene (including all isomers alone or in admixture).  
20 Preferred solvents are aliphatic hydrocarbons especially C<sub>5</sub>-C<sub>10</sub> alkanes or cycloalkanes and mixtures thereof. Such a mixture is available commercially under the trade designation Isopar™ E, available from Exxon Chemicals. If no cocatalyst is necessary, the complex itself is contacted with the addition polymerizable monomer.

"Addition polymerizable monomers" usefully polymerized according to the  
25 present invention include, for example, ethylenically unsaturated monomers, cyclic olefins, acetylenic compounds, conjugated or nonconjugated dienes and polyenes. Specific examples include C<sub>2-20</sub> olefins, styrene, halo- or hydrocarbyl substituted styrenes, divinylbenzene, 4-vinylcyclohexene, tetrafluoroethylene, vinylbenzocyclobutane, butadiene, isoprene, 1,4-hexadiene, cyclobutene, cyclopentene, cyclohexene, norbornene, ethylidene norbornene,  
30 acetylene, and mixtures thereof. Preferred monomers include the C<sub>2-10</sub> α-olefins especially ethylene, propylene, isobutylene, 1-butene, 1-hexene, 4-methyl-1-pentene, styrene, and 1-octene.

In general, the polymerization may be accomplished at conditions well known in the prior art for Ziegler-Natta or Kaminsky-Sinn type polymerization reactions, that is,  
35 temperatures from 0 to 250°C and pressures from atmospheric to 1000 atmospheres (100 MPa). Suspension, solution, slurry, gas phase or other process condition may be employed if desired. The catalyst may be supported and such supported catalyst may be employed in the polymerizations of this invention. Preferred supports include alumina and silica.

TABLE I

Ex.	Com- plex	Amt. ( $\mu$ mol)	Cocat- alyst	Amt. ( $\mu$ mol)	Solvent (g)	1-Octene (g)	Ethylene (MPa)	H <sub>2</sub> ( $\Delta$ kPa)	Temp (°C)	Time (min)	Polymer (g)	Efficiency (g poly/g Ti)
1	i	5.00	I	5.0	715	143	3.1	69	130	10	112.6	470,146
2	i	5.00	I	5.0	715	143	"	69	130	10	83.2	347,390
3	i	2.50	I	2.5	715	143	"	34	120	18	62.3	520,251
4	i	10.00	II	2940	715	143	"	340	130	15	54.7	114,196
5	i	10.00	II	184	715	143	"	340	130	"	22.5	46,973
6	i	10.00	II	2940	715	143	"	340	130	"	58.5	122,129
7	i	10.00	IV	1000	715	143	"	340	130	"	23.7	49,478
8	i	10.00	II	1470	787	72	"	34	160	"	17.8	37,161
9	i	10.00	II	2940	787	72	"	0	175	"	13.6	28,392
10	i	10.00	II	7350	715	143	"	340	130	"	53.7	112,109
11	iii	10.00	II	588	715	143	"	340	130	"	37.8	78,914
12	iii	10.00	II	441	715	143	"	340	130	"	23.1	48,225
13	i	0.50	I	1.5	740	118	3.4	170	100	"	62.2	2,597,077
14	i	1.50	I	3.0	788	70	"	90	160	"	20.0	285,317
"	i	1.00	I	3.0	740	118	"	340	140	"	30.7	640,919
16	i	1.00	I	3.0	740	118	"	0	140	"	18.7	390,397
17	i	0.75	I	1.5	691	167	"	90	120	"	45.5	1,266,527
18	i	0.75	I	1.5	788	70	"	260	120	"	44.8	1,247,042
19	i	0.75	I	3.0	691	167	"	90	120	"	43.2	1,202,505
20	i	1.00	I	3.0	740	118	"	170	140	"	17.2	359,081
21	i	1.50	I	6.0	788	70	"	260	160	"	12.4	172,582
22	i	0.75	I	1.5	788	70	"	90	120	"	41.6	1,157,968
23	i	1.00	I	3.0	740	118	"	170	140	"	10.1	210,856
24	i	0.75	I	3.0	788	70	"	90	120	"	45.2	1,258,177
25	i	1.00	I	3.0	740	118	"	170	140	"	16.3	340,292

TABLE I (continued)

Ex.	Com- plex	Amt. ( $\mu$ mol)	Cocat- alyst	Amt. ( $\mu$ mol)	Solvent (g)	1-Octene (g)	Ethylene (MPa)	H <sub>2</sub> ( $\Delta$ kPa)	Temp (°C)	Time (min)	Polymer (g)	Efficiency (g poly/g Ti)
26	i	1.00	I	3.0	740	118	3.4	170	140	15	12.6	263,048
27	i	10.00	I	30.0	740	118	"	"	180	"	12.5	26,096
28	i	1.00	I	3.0	837	21	"	"	140	"	43.0	897,704
29	i	1.00	I	3.0	740	118	"	"	140	"	27.9	582,463
30	i	0.75	I	3.0	691	167	"	260	120	"	39.4	1,096,729
31	i	1.00	I	3.0	740	118	"	170	140	"	20.2	421,712
32	i	1.50	I	6.0	691	167	"	91	160	"	5.1	70,981
33	i	1.50	I	3.0	691	167	"	91	160	"	5.8	80,724
34	i	1.00	I	3.0	644	215	"	170	140	"	13.8	288,100
35	i	1.50	I	3.0	691	167	"	260	160	"	8.4	116,910
36	i	1.00	I	5.0	740	118	"	170	140	"	17.0	354,906
37	i	1.50	I	6.0	691	167	"	260	160	"	6.9	96,033
38	i	0.75	I	1.5	691	167	"	260	120	"	50.9	1,416,841
39	i	1.50	I	3.0	788	70	"	260	160	"	7.3	101,601
40	i	0.75	I	3.0	788	70	"	260	120	"	29.7	826,722
41	i	1.00	I	1.0	740	118	"	170	140	"	7.8	162,839
42	i	1.50	I	6.0	788	70	"	90	160	"	2.9	40,362
43	i	1.00	I	3.0	740	118	"	170	140	"	13.3	277,662
44	i	2.50	III	2.5	740	118	"	"	120	"	1.6	13,361
45	i	2.50	III	5.0	740	118	"	"	120	"	.7	5,846
46	i	2.50	III	7.5	740	118	"	"	120	"	0.7	5,846
47	i	2.50	IV	7.5	740	118	"	"	120	"	0.5	4,175
48	i	2.50	V	37.5	740	118	"	"	120	"	3.3	20,577
49	i	2.50	VI	3.7	740	118	"	"	120	"	8.2	68,476
50	i	2.50	VI	7.5	740	118	"	"	120	"	8.9	74,322

TABLE I (continued)

Ex.	Com- plex	Amt. ( $\mu$ mol)	Cocat- alyst	Amt. ( $\mu$ mol)	Solvent (g)	1-Octene (g)	Ethylene (MPa)	H <sub>2</sub> ( $\Delta$ kPa)	Temp (°C)	Time (min)	Polymer (g)	Efficiency (g poly/g Ti)
51	i	2.50	VI	25.0	740	118	3.4	170	120	15	9.1	75,992
52	i	2.50	VI	75.0	740	118	"	"	"	"	5.7	47,599
53	iv	2.50	I	7.5	740	118	"	"	"	"	36.3	303,132
54	ii	2.00	I	2.0	740	118	"	"	140	"	112.4	1,174,000
55	ii	"	VII	3.0	"	"	"	"	140	"	122.4	1,277,000
56	ii	"	VIII	3.0	"	"	"	"	140	"	119.5	1,247,000
57	ii	"	-	0.0	"	"	"	"	"	"	72.3	755,000
58	v	0.75	I	0.75	708	150	"	"	120	"	68.8	1,915,000



Example 58

(i) Preparation of (phenylamido)dimethyl(tetramethyl- $\eta^5$ -cyclopentadienyl)silane 2-(N,N-dimethylamino)benzyl titanium chloride

In an argon filled glove box, 0.55 g (1.29 mmol) of (phenylamido)dimethyl-  
5 (tetramethyl- $\eta^5$ -cyclopentadienyl)silane titanium dichloride (prepared according to Example 89 of EP-A-0416845 and 0.18g (1.3 mmol) of 2-(N,N-dimethylamino)benzyl lithium were slurried in ~75 mL of pentane. The mixture turned from yellow to red after ~16 hours. The mixture was filtered and the solids were extracted with diethyl ether. The pentane and ether fractions were devolatilized under reduced pressure. <sup>1</sup>H NMR analysis confirmed the product's identity from  
10 both fractions as (phenylamido)dimethyl(tetramethyl- $\eta^5$ -cyclopentadienyl)silane titanium 2-(N,N-dimethylamino)benzyl chloride. Total yield was 0.33 g, 53 percent.

(ii) Preparation of (phenylamido)dimethyl(tetramethyl- $\eta^5$ -cyclopentadienyl)silane-2-(N,N-dimethylamino)benzyl titanium

In an argon filled glove box 16 mg of magnesium powder was added to 0.33 g  
15 (0.68 mmol) of (phenylamido)dimethyl(tetramethyl- $\eta^5$ -cyclopentadienyl)silane 2-(dimethylamino)benzyl titanium chloride in ~40 mL of tetrahydrofuran (THF). After stirring for 6 hours the THF was removed under reduced pressure. The solid residue was extracted with pentane. The pentane extracts were filtered and combined and the pentane solvent removed under  
20 reduced pressure to give 0.29 g (94 percent yield) of (phenylamido)dimethyl(tetramethyl- $\eta^5$ -cyclopentadienyl)silane-2-(N,N-dimethylamino)benzyl titanium, characterized by cyclic voltametry.

Example 59

25 A 100 mil round bottom flask was loaded with 0.100 g of (tert-butyl-amido)dimethyl(tetramethyl- $\eta^5$ -cyclopentadienyl)silane 2-(N, N-dimethylamino)benzyl titanium. 30 ml of pentane were added to form a solution. To this solution was added, via pipet, 0.119 g of B(C<sub>6</sub>F<sub>5</sub>)<sub>3</sub> in 40 ml of pentane. Within minutes the solution become cloudy and a pink/orange solid precipitated from solution. The reaction mixture was stirred for an  
30 additional 15 minutes after which time the solid was collected by filtration and washed with fresh pentane (3x15 ml). Yield of the product (believed to be [(tert-butyl-amido)dimethyl(tetramethyl- $\eta^5$ -cyclopentadienyl)silane titanium(III)] [(2-N, N-dimethylamino)benzyl)trispentafluorophenylboron), after drying under reduced pressure was 0.205 g.

35 The ESR spectrum of this reaction product showed a single line at room temperature (g = 1.97). The cycle voltamagram (in dimethyl ether (DME) with no supporting electrolyte) of the reaction product showed an irreversible reduction wave at -2.74 volts vs. ferrocene. Under similar conditions, the neutral complex, (tert-butyl-

amido)dimethyl(tetramethyl- $\eta^5$ -cyclopentadienyl)silane (2-N, N-dimethylamino)benzyl titanium, was not conductive and did not give a cyclic voltamogram. The fact that electrochemical measurements could be obtained in the absence of added supporting electrolyte strongly indicates that the activated complex was cationic.

- 5 Under the polymerization conditions of Example 54, the above activated complex used as a catalyst gave 28.8 g of ethylene/octene copolymer.

#### Example 60

- 10 The reaction conditions of Example 59 were repeated to prepare the pink/orange cationic metal complex which was then redissolved in tetrahydrofuran thereby giving a purple solution. Addition of pentane caused precipitation of a purple solid which was collected by filtration, washed with fresh pentane, and dried.

- The ESR spectrum of the purple product showed a single line at room temperature ( $g = 1.97$ ) and the cyclic voltamogram (in DME with no supporting electrolyte) showed an irreversible reduction wave at -2.74 volts vs. ferrocene. Under similar conditions, the neutral complex did not give a cyclic voltamogram thereby indicating that the purple solid was an ionic  $Ti + 3$  complex.

- 15 15 mg of the purple solid was combined with 0.5 mL of methanol and the resulting solution was analyzed by GC/mass spectroscopy. THF was identified in the solution, indicating that the purple solid contained the neutral Lewis base, THF. On this basis the purple solid was believed to be [(tert-butylamido)dimethyl(tetramethyl- $\eta^5$ -cyclopentadienyl)silane titanium(III)(THF)] ((2-N, N-dimethylamino)benzyl)tris(pentafluorophenyl)boron).

- 25 Under the polymerization conditions of Example 54, the above purple solid used as a catalyst gave 29.2 g of ethylene/octene copolymer.

#### Example 61

- (i) Preparation of (t-butylamido)dimethyl(tetramethyl- $\eta^5$ -cyclopentadienyl)silane (1-phenyl-3-benzylallyl) titanium

- 30 In an argon filled glove box, 0.50 g (1.4 mmol) of (t-butylamido)dimethyl-(tetramethyl- $\eta^5$ -cyclopentadienyl)silane titanium dichloride was dissolved into 30 mL of diethyl ether to give a yellow solution. 0.28 g (1.4 mmol) of 1,4-diphenyl-1,3 butadiene and 1.18 mL (2.8 mmole) of isopropyl magnesium chloride were added with stirring. Gas evolution occurred and the mixture darkened to a red/purple color over 50 minutes. The solvent was removed and the product was dried. The solid was extracted with pentane (5x10 mL) and dried under reduced pressure to give the product as a red/purple solid (0.545 g).
- 35

CLAIMS

1. An addition polymerization catalyst comprising:

(i) a metal complex corresponding to the formula:  $\text{Cp}_a(\text{ZY})_b\text{ML}_c$ , wherein:

a is 1 or 2;

5 b is 0 or 1;

c is 1 or 2;

the sum of a, b and c is 3;

Cp independently each occurrence is a cyclopentadienyl group  $\eta$ -bound to M, or a hydrocarbyl, silyl, halo, halohydrocarbyl, hydrocarbylmetaloid or halohydrocarbylmetaloid substituted derivative of said cyclopentadienyl group, said Cp containing up to 50 nonhydrogen atoms, and, when a is 2, optionally both Cp groups may be joined together by a bridging group;

L independently each occurrence is hydride, halo, or a monovalent anionic ligand selected from covalently bonded hydrocarbyl, silyl, amido, phosphido, alkoxy, aryloxy, and sulfido groups optionally being further substituted with amine, phosphine, ether, and thioether; mixtures thereof; said ligand having up to 50 nonhydrogen atoms, with the proviso that in at least one occurrence L is a stabilizing ligand comprising an amine, phosphine, ether or thioether functionality able to form a coordinate-covalent bond or chelating bond with M, or except when a is 1, comprising an ethylenic unsaturation able to form an  $\eta^3$  bond with M;

20 M is a metal of Group 4 of the Periodic Table of the Elements in the + 3 oxidation state;

Y is a linking group comprising nitrogen, phosphorus, oxygen or sulfur covalently bonded to M and Z through said nitrogen, phosphorus, oxygen or sulfur atom;

25 Z is a divalent moiety comprising oxygen, nitrogen, phosphorous, boron, or a member of Group 14 of the Periodic Table of the Elements, said moiety having up to 30 nonhydrogen atoms;

the ligand moiety consisting of -Cp-Z-Y- being dianionic and having the ionic charges residing formally on Cp and Y; and, when b is 0 and optionally when b is 1; and (ii) an activating cocatalyst.

30 2. An addition polymerization catalyst comprising in combination:

(i) a metal complex corresponding to the formula:  $\text{Cp}'_2\text{ML}'$ , wherein:

Cp' independently each occurrence is a cyclopentadienyl group  $\eta$ -bound to M, or a hydrocarbyl, silyl, halo, halohydrocarbyl, hydrocarbylmetaloid or halohydrocarbylmetaloid substituted derivative of said cyclopentadienyl group, said Cp' containing up to 50 nonhydrogen atoms, and optionally both Cp groups may be joined together by a bridging group;

M is a metal of Group 4 of the Periodic Table of the Elements in the +3 oxidation state;

L' is a monovalent anionic stabilizing ligand selected from covalently bonded hydrocarbyl, silyl, amido, phosphido, alkoxy, aryloxy, sulfido groups and mixtures thereof, and being further substituted with an amine, phosphine, ether, or thioether containing substituent able to form a coordinate-covalent bond or chelating bond with M; said ligand having up to 50 nonhydrogen atoms; and

(ii) an activating cocatalyst.

3. An addition polymerization catalyst comprising in combination:

(i) a metal complex corresponding to the formula:  $\text{Cp}''\text{ML}''_2$ , wherein:

$\text{Cp}''$  is a cyclopentadienyl group  $\eta$ -bound to M, or a hydrocarbyl, silyl, halo, halohydrocarbyl, hydrocarbylmetaloid, or halohydrocarbylmetaloid substituted derivative thereof, said  $\text{Cp}''$  containing up to 50 nonhydrogen atoms;

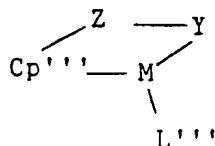
M is a metal of Group 4 of the Periodic Table of the Elements in the +3 oxidation state;

L'' independently each occurrence is hydride, halo, or a monovalent anionic ligand selected from covalently bonded hydrocarbyl, silyl, amido, phosphido, alkoxy, aryloxy, and sulfido groups; mixtures thereof; and amine, phosphine, ether, and thioether derivatives of the foregoing, said ligand having up to 50 nonhydrogen atoms, with the proviso that in at least one occurrence L'' is a stabilizing ligand comprising an amine, phosphine, ether or thioether functionality able to form a coordinate-covalent bond or chelating bond with M, or comprising an ethylenic unsaturation able to form an  $\eta^3$  bond with M; and

(ii) an activating cocatalyst.

25

4. A metal complex corresponding to the formula:



wherein:

M is a metal of Group 4 of the Periodic Table of the Elements in the +3 oxidation state;

$\text{Cp}'''$  is a cyclopentadienyl group, or a hydrocarbyl, silyl, halo, halohydrocarbyl, hydrocarbylmetaloid, or halohydrocarbylmetaloid substituted derivative thereof, said  $\text{Cp}'''$  containing up to 50 nonhydrogen atoms;

Z is a divalent moiety comprising oxygen, nitrogen, phosphorous, boron, or a member of Group 14 of the Periodic Table of the Elements, said moiety having up to 30 nonhydrogen atoms;

Y is a linking group comprising nitrogen, phosphorus, oxygen or sulfur covalently bonded to M and Z through said nitrogen, phosphorus, oxygen or sulfur atom, the ligand moiety consisting of  $\text{-Cp}'''\text{-Z-Y-}$  being dianionic and having the ionic charges residing formally on  $\text{Cp}'''$  and Y; and

$\text{L}'''$  is a monovalent anionic stabilizing ligand selected from:

- a) covalently bonded hydrocarbyl, silyl, amido, phosphido, alkoxy, aryloxy, sulfido groups and mixtures thereof, and being further substituted with an amine, phosphine, ether, or thioether containing substituent able to form a coordinate-covalent bond or chelating bond with M, said ligand having up to 50 nonhydrogen atoms; and
- b) linear  $\text{C}_3\text{-C}_{15}$  hydrocarbyl groups comprising an ethylenic unsaturation able to form an  $\eta^3$  bond with M.

15

5. An addition polymerization catalyst comprising a complex as claimed in Claim 4 and an activating cocatalyst.

6. A catalyst or complex as claimed in any one of the preceding claims, wherein M is titanium.

7. A catalyst or complex as claimed in any one of the preceding claims, wherein L, L', L'' or L''' is 2-di( $\text{C}_1\text{-C}_4\text{-alkyl}$ )aminobenzyl or 2-(di( $\text{C}_1\text{-C}_4\text{-alkyl}$ )aminomethyl)phenyl.

8. A catalyst or complex as claimed in Claim 7, wherein L, L', L'' or L''' is 2-(N,N-dimethylamino)benzyl.

9. A catalyst or complex as claimed in any one of Claims 1, and 3 to 6, wherein L, L'' or L''' is linear  $\text{C}_3\text{-C}_{15}$  hydrocarbyl comprising an ethylenic unsaturation.

30

10. A catalyst or complex as claimed in Claim 9, wherein L, L'' or L''' is allyl.

11. A catalyst as claimed in any one of Claims 1, 2, 6, 7 or 8, wherein the metal complex comprises two Cp groups joined by a bridging group of the formula  $(\text{R}''_2\text{E})_x$  wherein E is silicon or carbon and  $\text{R}''$  independently each occurrence is hydrogen, silyl, hydrocarbyl or a combination of silyl and hydrocarbyl, said  $\text{R}''$  having up to 30 silicon or carbon atoms, and x is 1 to 8.

35

12. A catalyst or complex as claimed in Claim 8, wherein the metal complex is biscyclopentadienyl 2-(N,N-dimethylamino)benzyl titanium (III).

13. A complex as claimed in Claim 4 selected from (tert-butylamido)-  
5 dimethyl(tetramethyl- $\eta$ 5-cyclopentadienyl)silane 2-(N,N-dimethylamino)benzyl titanium (III),  
(tert-butyl- amido)dimethyl(tetramethyl- $\eta$ 5-cyclopentadienyl)silane allyl titanium (III), (tert-  
butyl- amido)dimethyl(tetramethyl- $\eta$ 5-cyclopentadienyl)silane 1,2,3-trimethylallyl titanium  
(III), (tert-butyl- amido)dimethyl(tetramethyl- $\eta$ 5-cyclopentadienyl)silane 1-phenyl-3-benzylallyl  
titanium (III), and (tert-butyl- amido)dimethyl(tetramethyl- $\eta$ 5-cyclopentadienyl)silane 1,1-  
10 diphenyl-3-(diphenylmethyl)allyl titanium (III)

14. A catalyst as claimed in Claim 5, wherein the metal complex is selected from  
(tert-butylamido)-dimethyl(tetramethyl- $\eta$ 5-cyclopentadienyl)-silane 2-(N,N-  
dimethylamino)benzyl titanium (III), (tert-butylamido)dimethyl(tetramethyl- $\eta$ 5-  
15 cyclopentadienyl)silane allyl titanium (III), and (tert-butylamido)dimethyl(tetramethyl- $\eta$ 5-  
cyclopentadienyl)silane 1,2,3-trimethylallyl titanium (III).

15. An addition polymerization catalyst as claimed in Claim 12 or 14, wherein  
the cocatalyst is tris(pentafluorophenyl)boron.

20

16. A catalyst as claimed in any one of Claims 1 to 3, 5 to 12, 14 or 15, wherein  
the ratio of metal complex to activating cocatalyst is from 1:0.01 to 1:10<sup>6</sup>.

17. A catalyst as claimed in any one of Claims 1 to 3, 5 to 12 or 14 to 16,  
25 wherein the activating cocatalyst is selected from alkylalumoxanes; aluminum alkyls;  
aluminum halides; alkylaluminum halides; Lewis acids other than any of the foregoing; and  
mixtures thereof.

18. A catalyst as claimed in Claim 17, wherein the activating cocatalyst is tris(4-  
30 fluorophenyl)boron, tris(2,4-difluorophenyl)boron, tris(3,5-bis(trifluoromethyl)phenyl)boron,  
tris(pentafluorophenyl)boron, pentafluorophenyl diphenylboron, triphenyl boron or  
bis(pentafluorophenyl)phenylboron.

19. A catalyst as claimed in Claim 18, wherein the activating cocatalyst is  
35 tris(pentafluorophenyl)boron.

20. A catalyst or complex as claimed in any one of the preceding claims,  
additionally comprising a neutral Lewis base.

21. A catalyst or complex as claimed in claim 20, wherein the neutral Lewis base is selected from the group consisting of organic amines, amides, ketones, nitriles, phosphines, phosphorylamides, esters, ethers, thioethers and thioesters, said neutral Lewis  
5 base containing up to 20 nonhydrogen atoms.

22. A catalyzed addition polymerization process comprising contacting under addition polymerization conditions at least one addition polymerizable monomer with a catalyst or complex according to any one of Claims 1 to 21.

10 23. A process for preparing a catalyst as claimed in Claim 1, comprising contacting a Group 4 metal complex corresponding to the formula:  $Cp_a(ZY)_bM^*L_cX$ , wherein:

a, b, c, Cp, L, Y and Z are as defined in Claim 1,  
15  $M^*$  is a metal of Group 4 of the Periodic Table of the Elements in the + 4 oxidation state; and

X is halide or  $C_{1-10}$  hydrocarbyloxyde,  
with a reducing agent under reducing conditions to form a metal complex corresponding to the formula:  $Cp_a(ZY)_bML_c$ ,  
20 wherein a, b, c, Cp, L, M, Y and Z are as defined in Claim 1,  
and, when a cocatalyst is present, contacting the so formed complex with the cocatalyst.

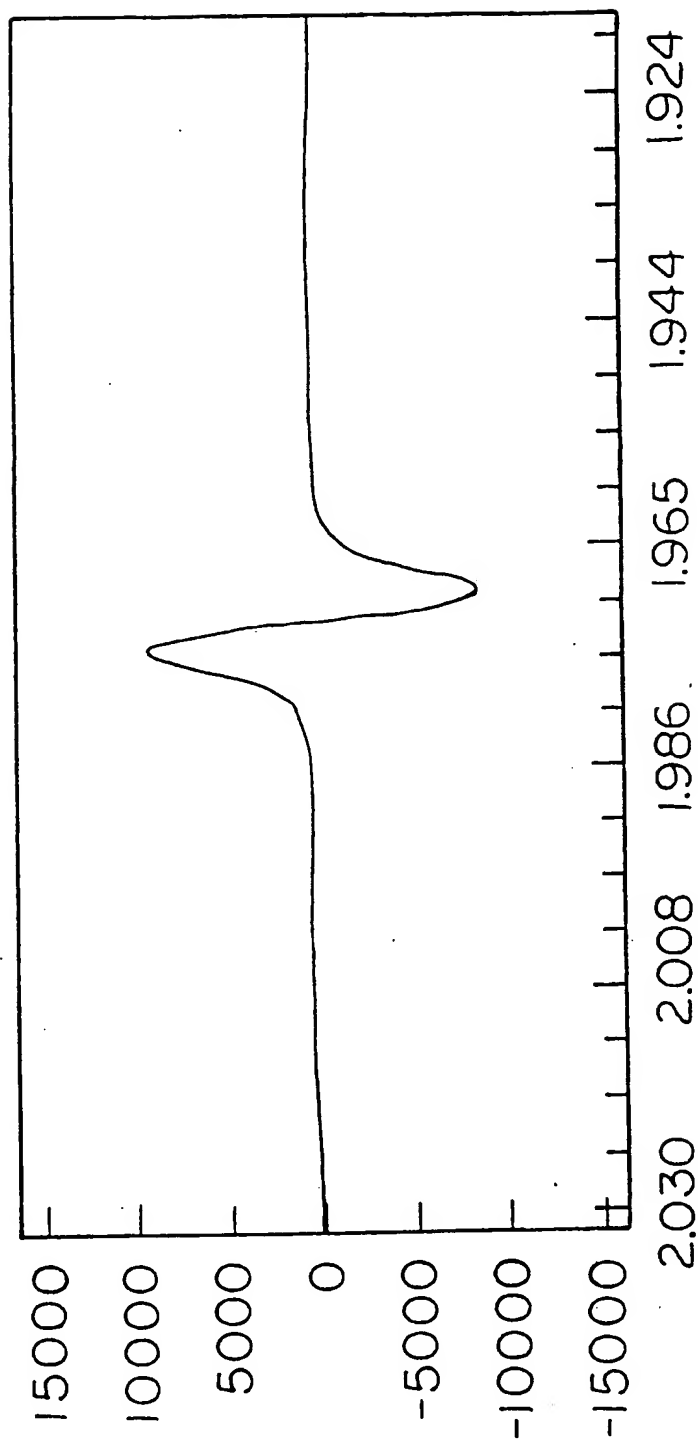
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FIG. 1

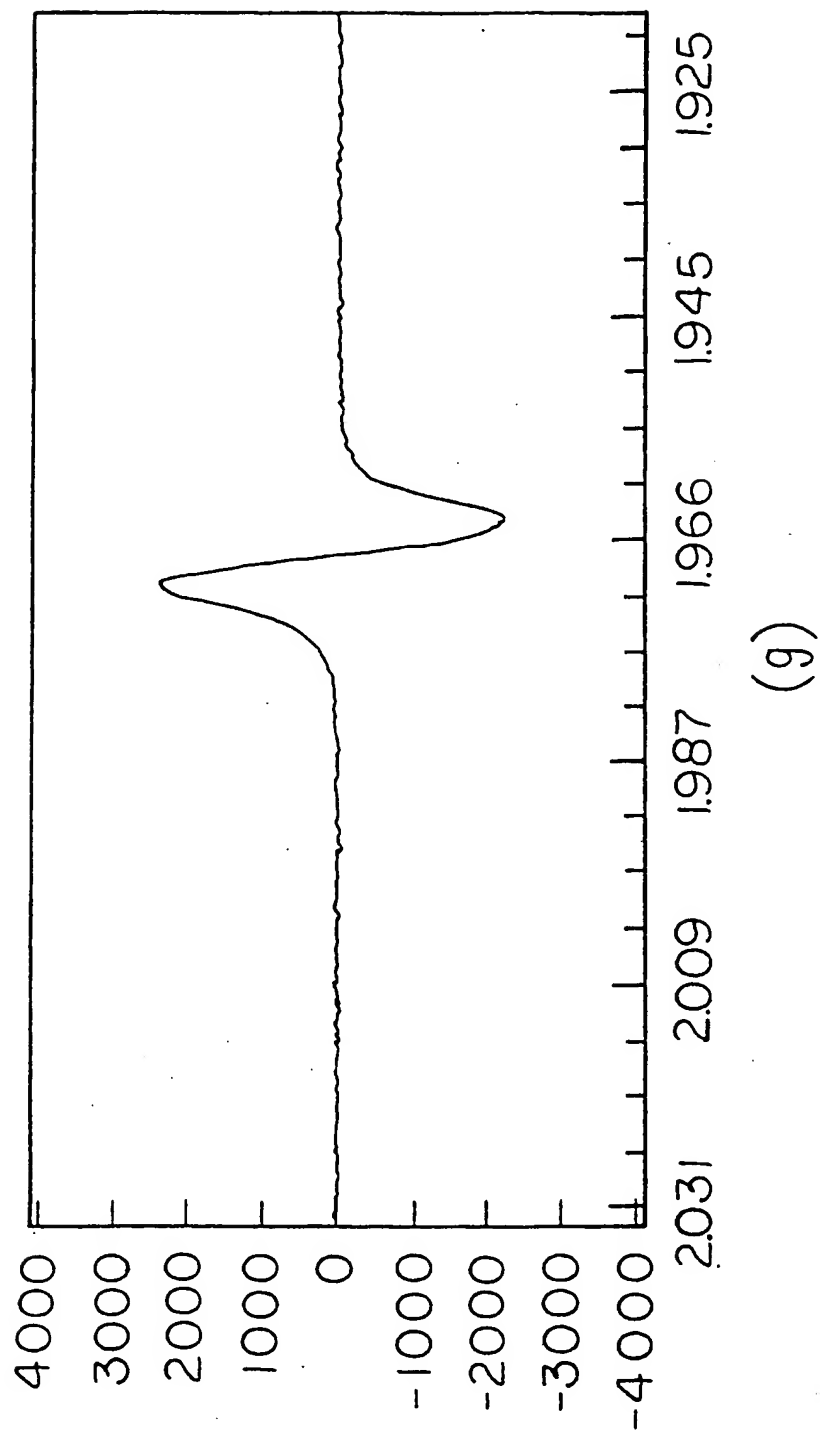


(g)



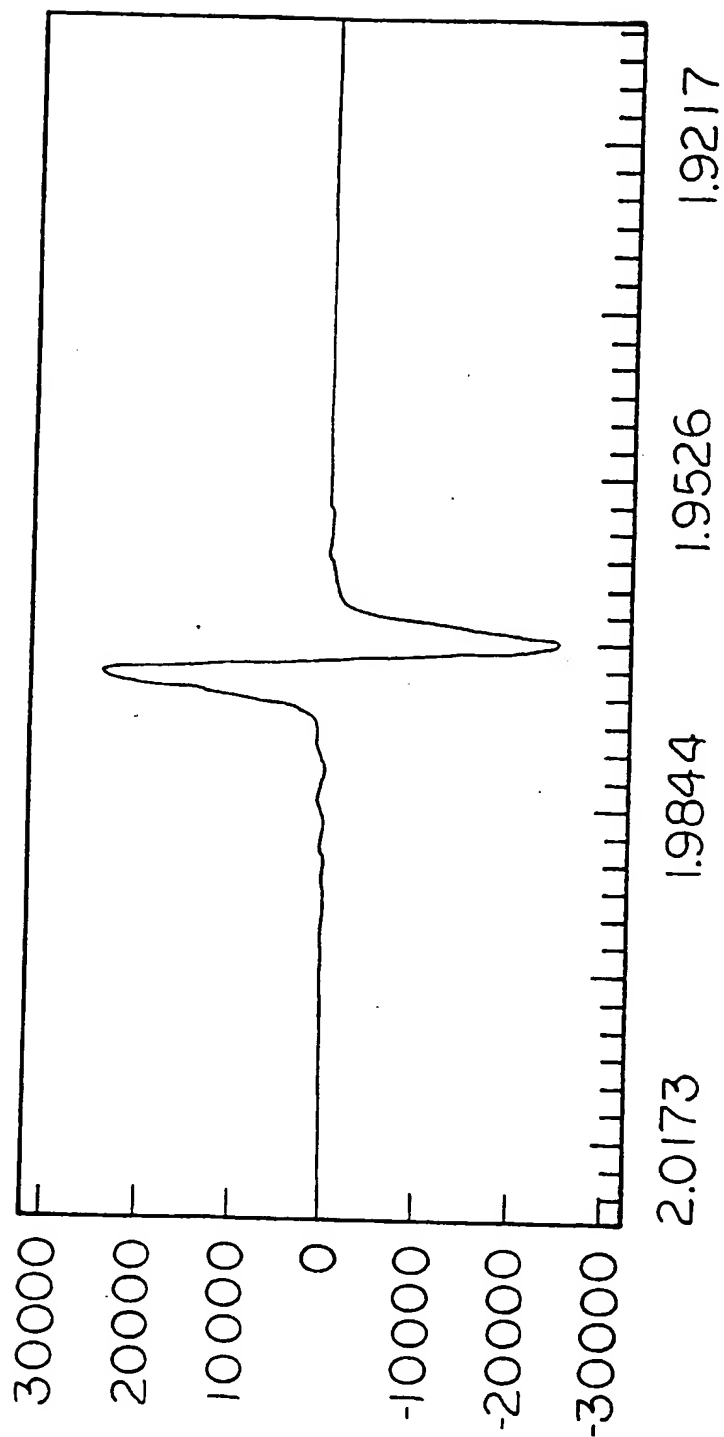
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FIG. 2



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FIG. 3



ANNEX TO THE INTERNATIONAL SEARCH REPORT  
ON INTERNATIONAL PATENT APPLICATION NO.

US 9302584  
SA 71723

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Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP-A-0416815	13-03-91	AU-A- 6203990	07-03-91
		CA-A- 2024333	01-03-91
		JP-A- 3163088	15-07-91
		CN-A- 1049849	13-03-91
EP-A-0495375	22-07-92	US-A- 5189192	23-02-93
		AU-A- 1023392	23-07-92
		JP-A- 5086120	06-04-93
EP-A-0468537	29-01-92	AU-B- 617990	12-12-91
		AU-A- 1245288	24-08-88
		EP-A- 0277004	03-08-88
		JP-T- 1502036	13-07-89
		WO-A- 8805793	11-08-88
		US-A- 5153157	06-10-92
		US-A- 5198401	30-03-93

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